#### 4.0 MEASURED CONCENTRATION TRENDS

### 4.1 TRENDS OF MEASURED SO<sub>2</sub> CONCENTRATIONS

Monitors for SO<sub>2</sub> have been maintained in the Theodore Roosevelt National Park (TRNP), South Unit and North Unit, since the early 1980s. This covers virtually all of the period since the PSD baseline date was established for the North Dakota PSD Class I areas. If it were true that the PSD increment has been totally consumed, then this feature should show up in the monitoring data as an increase in the second-highest concentrations over time to the present day.

Figures 4-1 and 4-2 show the trends of the 3-hour and 24-hour highest, second highest measurements at the TRNP North Unit, while Figures 4-3 and 4-4 provide the trends at the TRNP South Unit monitor. It is clear from the figures that the recent SO<sub>2</sub> measurements are among the lowest in the past 20 years. This is evidence that the air quality in the PSD Class I areas associated with the highest modeled results has not degraded, and has actually improved. Therefore, the model emission inventory must be missing some key PSD increment expanding sources that would lead to this result. More discussion of this issue is provided in Section 4-2.

#### 4.2 PSD EMISSIONS INVENTORY

The EPA report states that the air quality in the PSD Class I areas is only slightly affected by the nearby oil and gas sources. However, the proximity of these sources to the Class I areas creates considerable doubt as to the validity of that assumption, particularly in light of the monitoring trends shown in Section 4.1. If the observed concentrations of SO<sub>2</sub> are not increasing in the PSD Class I areas (an observation which has led to permit variances granted in the past), then it does not make sense that modeling results show an increment violation. The CALPUFF predictions within each Class I area from the distant sources are relatively uniform, given the small sizes of the parks in relation to the distances involved from the major SO<sub>2</sub> sources. The concentration trend is consistent between the TRNP North and South Unit monitors. The only explanation for this inconsistency is that some PSD increment expansion sources are not being accounted for in the modeling.

Where are these PSD increment expansion sources, and when did they operate? The answer may lie with the nearby oil and gas producing sources that EPA has not yet considered. It is likely that in the 1970s, the lack of available gas pipelines caused these sources to continuously flare gas streams that could not otherwise be marketed. Later on, the construction of gas pipelines allowed the gas streams to be marketed, and the flaring operations closed down. These flare emissions, plus other emissions associated with the numerous nearby oil and gas sources, should be accounted for in the modeling as accurately and completely as possible.

Another important increment expanding source is the Royal Oak briquette factory near Dickinson. This source was only about 50 kilometers from the TRNP South Unit and it was a major  $SO_2$  source. It is noteworthy that in its updated assessment of baseline source emissions, the NDDH has more than tripled the  $SO_2$  emission rate from Royal Oak, from about 69 to 222 grams per second. It is also noteworthy that during the maximum coal usage period of this facility (during the 1980s), the observed  $SO_2$  concentrations at the TRNP South and North Unit monitors registered their highest concentrations, as shown in Figures 4-1 through 4-4. The coincidence of these emissions and the corresponding monitored peaks is worthy of further investigation.

In summary, EPA needs to more thoroughly review the baseline emissions so that they can account for the overall decrease in measured  $SO_2$  concentrations over the past 20+ years. Until this step is accomplished, the EPA study cannot be considered as being satisfactorily completed or credible.

4-2



Figure 4-1 Monitored SO<sub>2</sub> Values – TRNP-NU 3-hour 2<sup>nd</sup> High

## Monitored SO2 Values - TRNP-NU 3-hour 2nd High



Figure 4-2 Monitored SO<sub>2</sub> Values – TRNP-NU 24-hour 2<sup>nd</sup> High

# Monitored SO2 Values - TRNP - NU 24-hour 2nd High

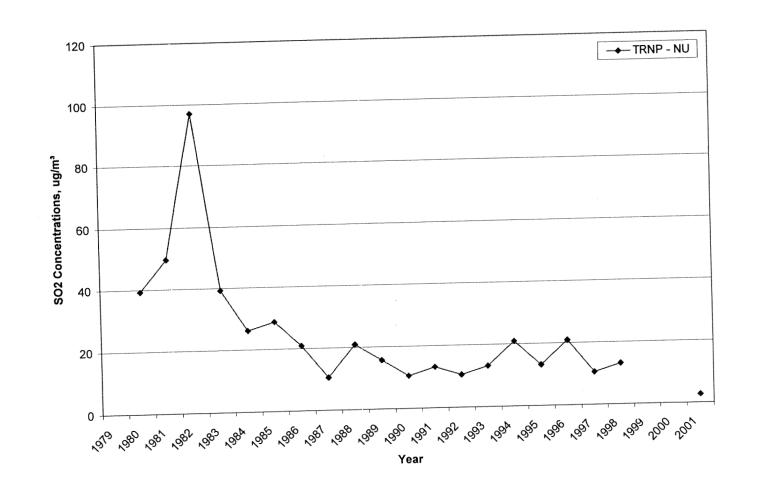


Figure 4-3 Monitored SO<sub>2</sub> Values – TRNP-SU 3-hour 2<sup>nd</sup> High

## Monitored SO2 Values - TRNP - SU 3-hour 2nd High

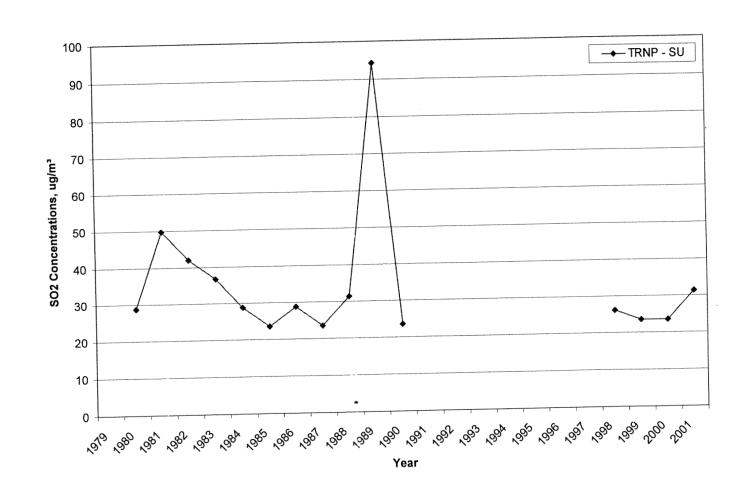
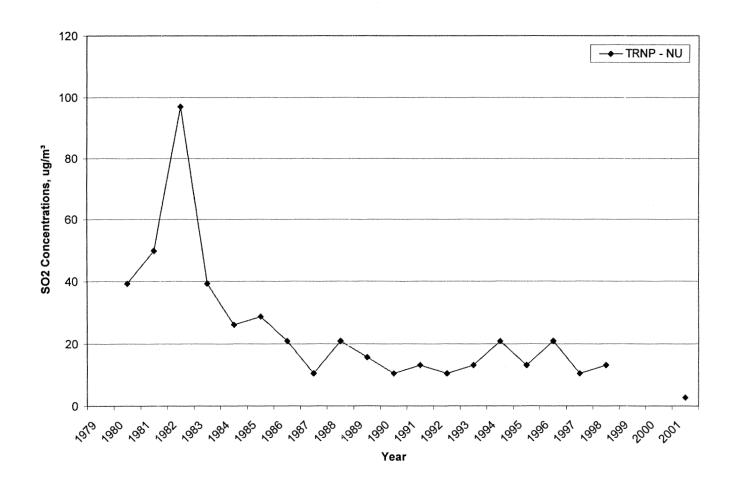




Figure 4-4 Monitored SO<sub>2</sub> Values – TRNP-NU 24-hour 2<sup>nd</sup> High

### Monitored SO2 Values - TRNP - NU 24-hour 2nd High





### 5.0 CONCLUSIONS

The major comments that are provided in this document are summarized below.

- While the CALPUFF model is an advancement over previous techniques for long-range transport modeling, the model still has significant limitations. At the distances being considered between the major sources and PSD Class I receptors, CALPUFF would be expected to over predict by about a factor of 2, based upon results from independent studies.
- The NDDH CALPUFF evaluation for the year 2000 neglected to consider regional background concentration contributions to the full predicted concentration. When a low regional background value of 4 μg/m3 is accounted for, the evaluation results show an over prediction tendency of about a factor of 2, in agreement with the studies reported by IWAQM.
- Monitoring data in the TRNP North and South Units provides evidence that the SO<sub>2</sub> concentrations have, if anything, dropped over the past 20 years. The modeling should indicate a similar trend. Since it does not, the only explanation is that some PSD increment expanding sources have not yet been accounted for. Possible baseline emission contributors to the past high observed SO<sub>2</sub> concentrations are flares at numerous oil and gas wells and the Royal Oak briquette plant. Since these past sources were much closer to the affected PSD Class I areas than most of the major SO<sub>2</sub> increment consuming sources assessed in this study, these emission reductions could account for the observed improvement in air quality within the PSD Class I areas.

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